	Set Name side by side	Query	Hit Count	Set Name result set
	DB=USPT,F	PGPB,JPAB,EPAB,DWPI,TDBD; PLUR=YES; OP=ADJ	•	
	<u>L14</u>	polyacetal-polyether	2	<u>L14</u>
•	<u>L13</u>	polyacetal near1 polyether near1511	2	<u>L13</u>
	<u>L12</u>	polyacetal near2 polyether near1511	10	<u>L12</u>
	<u>L11</u>	polyacetal near3 polyether near1511	15	<u>L11</u>
	<u>L10</u>	polyacetal near5 polyether near1511	19	<u>L10</u>
,	<u>L9</u>	L5 near15 11	0	<u>L9</u>
	<u>L8</u>	L5 near5 11	0	<u>L8</u>
	<u>L7</u>	L5 near 11	0	<u>L7</u>
	<u>L6</u>	L5 and 11	13	<u>L6</u>
	<u>L5</u>	thickener	49236	<u>L5</u>
	<u>L4</u>	polyacetal near polyether [clm]	42	<u>L4</u>
	<u>L3</u>	polyacetal near polyether [ab]	73	<u>L3</u>
	<u>L2</u>	polyacetal near polyether [ti]	4	<u>L2</u>
	<u>L1</u>	polyacetal near polyether	575	<u>L1</u>
	6.5			

END OF SEARCH HISTORY

10/003755

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Search Results - Record(s) 1 through 3 of 3 returned.

1. Document ID: US 20020006985 A1

L6: Entry 1 of 3

File: PGPB

Jan 17, 2002

PGPUB-DOCUMENT-NUMBER: 20020006985

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020006985 A1

TITLE: FINE PARTICLE SIZE LOW BULK DENSITY THERMOPLASTIC POLYMERS

PUBLICATION-DATE: January 17, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

SAU, ARJUN CHANDRA

NEWARK

DE

US

ТОПЕ

US-CL-CURRENT: 523/332; 524/503, 525/398, 525/924 .

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KWC | Draw, Desc | Image |

2. Document ID: US 6369132 B2

L6: Entry 2 of 3

File: USPT

Apr 9, 2002

US-PAT-NO: 6369132

DOCUMENT-IDENTIFIER: US 6369132 B2

TITLE: Fine particle size low bulk density thermoplastic polymers

DATE-ISSUED: April 9, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Sau; Arjun Chandra

Newark

DE

US-CL-CURRENT: 523/332; 523/340, 524/549, 524/556, 524/559, 524/592, 524/593, 524/612, 525/398, 525/471, 525/92K, 528/501, 536/43, 536/44

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KMC Draw. Desc Image

3. Document ID: US 3624012 A

L6: Entry 3 of 3

File: USPT

Nov 30, 1971

US-PAT-NO: 3624012

DOCUMENT-IDENTIFIER: US 3624012 A

TITLE: PROCESS FOR PREPARING HALOGENATED COPOLYMERS OF OLEFINICALLY UNSATURATED

CYCLOALIPHATIC MONOANHYDRIDES WITH CYCLIC MONOETHERS

DATE-ISSUED: November 30, 1971

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Case; Leslie C.

Winchester

MA

01890

US-CL-CURRENT: 524/724; 524/701, 524/706, 524/732, 524/734, 524/736, 524/738, 524/739, 524/750, 524/761, 528/366

524/750, 524/761, 528/366

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments
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Term	Documents
POLYETHER.DWPI,TDBD,EPAB,JPAB,USPT,PGPB.	89953
POLYETHERS.DWPI,TDBD,EPAB,JPAB,USPT,PGPB.	25564
POLYACETAL.DWPI,TDBD,EPAB,JPAB,USPT,PGPB.	14753
POLYACETALS.DWPI,TDBD,EPAB,JPAB,USPT,PGPB.	7984
CYCLODEXTRIN.DWPI,TDBD,EPAB,JPAB,USPT,PGPB.	13184
CYCLODEXTRINS.DWPI,TDBD,EPAB,JPAB,USPT,PGPB.	4493
((POLYACETAL NEAR POLYETHER) AND CYCLODEXTRIN).USPT,PGPB,JPAB,EPAB,DWPI,TDBD.	3
(POLYETHER NEAR POLYACETAL AND CYCLODEXTRIN).USPT,PGPB,JPAB,EPAB,DWPI,TDBD.	3

Display Format: -

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L6: Entry 3 of 3

File: USPT

Nov 30, 1971

DOCUMENT-IDENTIFIER: US 3624012 A

TITLE: PROCESS FOR PREPARING HALOGENATED COPOLYMERS OF OLEFINICALLY UNSATURATED CYCLOALIPHATIC MONOANHYDRIDES WITH CYCLIC MONOETHERS

Brief Summary Text (5):

This invention relates to the preparation of halogen-containing binary and ternary copolymers related to the classes broadly recognized as polyester, polyether and polyacetal compositions. More specifically, this invention is concerned with an improved two-step process for preparing those copolymers which possess, incorporated into the polymer chain, monomer residues derived from halogenated cycloaliphatic dicarboxylic acid monoanhydrides.

Brief Summary Text (16):

I have now discovered that the aforementioned disadvantages are completely eliminated by adding halogen not to the ethylenically unsaturated cycloaliphatic dicarboxylic anhydride but to the unsaturated copolymer prepared therefrom. Quite surprisingly, I have found that the addition of halogen to such an unsaturated copolymer proceeds smoothly and without cleavage of, or other undesirable interaction with, the polyether and polyacetal segments of the copolymer chains. The post-halogenation process of the present invention avoids any loss of raw materials due to side reactions and resinification, eliminates the disadvantages associated with the preparation of the halogenated monomer and results in polymers of good color.

Detailed Description Text (25):

Preferred for use are monosaccharides, disaccharides and trisaccharides having from five to 18 carbon atoms and from four to 11 hydroxyl groups. Especially preferred are monosaccharides and disaccharides. Examples of especially suitable compounds include among others dextrose, also known as glucose, xylose, sucrose, lactose, and maltose. Higher polysaccharides having more than three monosaccharide unit building blocks per molecule are also suitable. Examples of such higher polysaccharides are linear, branched and cyclic dextrins, such as alpha-cyclodextrin and beta-cyclodextrin, and polysaccharides of less well-defined composition, such as molasses, plant gums, mucilages, dextrans, pectins, corn syrup, and solubilized starches, such as solubilized corn starch, or potato starch, and the like. Products resulting from the modification and degradation of cellulose, such as partially esterified cellulose acetate may also be of value.

<u>Detailed Description Text</u> (27):

Particularly preferred for use are the white corn dextrins, the canary corn dextrins, British Gums, cyclodextrins, corn starch, potato starch, and wheat starch. The proportion of polysaccharide in the polysaccharide-polyol starter mixture will generally range from about 5 percent to about 90 percent, and advisably to no more than 80 percent, and preferably from about 20 percent to about 60 percent by weight based on the weight of the combined starter mixture. These polyol-polysaccharide mixtures provide polymerization starters of extraordinarily high functionality, since the hydroxyl functionality of the polysaccharide component of the starter mixture may range from about 10 up to about 100,000, or even more.